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**UTILITY
PATENT APPLICATION
TRANSMITTAL**

(Only for new nonprovisional applications under 37 CFR 1.53(b))

Attorney Docket No. 2000-1578A

Total Pages :

First Named Inventor or Application Identifier

Yoichi MORI et al.

Express Mail Label No.:

APPLICATION ELEMENTS

See MPEP chapter 600 concerning utility patent application contents.

ADDRESS TO: Assistant Commissioner for Patents
Box Patent Application
Washington, D.C. 20231

1. ☒ Fee Transmittal Form
(Submit an original, and a duplicate for fee processing)

6. Microfiche Computer Program (Appendix)

2. ☒ Specification [Total Pages - 13]
(preferred arrangement set forth below)
- Descriptive title of the Invention
- Cross References to Related Applications
- Statement Regarding Fed sponsored R & D

7. ☐ Nucleotide and/or Amino Acid Sequence Submission
(if applicable, all necessary)

- a. ☐ Computer Readable Copy
b. ☐ Paper Copy (identical to computer copy)
c. ☐ Statement verifying identity of above copies

- Reference to Microfiche Appendix
- Background of the Invention
- Brief Summary of the Invention
- Brief Description of the Drawings (if filed)
- Detailed Description
- Claim(s)
- Abstract of the Disclosure

ACCOMPANYING APPLICATION PARTS

3. ☒ Drawing(s) (35 USC 113) [Total Sheets - 1]

8. ☒ Assignment Papers (cover sheet & document(s))

4. ☒ Oath or Declaration [Total Pages - 3]

9. ☐ 37 CFR 3.73(b) Statement ☐ Power of Attorney
(when there is an assignee)

a.1. ☒ Newly executed (original or copy)

10. ☐ English Translation Document (if applicable)

a.2. ☐ Unexecuted

11. ☐ Information Disclosure Statement (IDS)/PTO-1449
☐ Copies of IDS Citations

b. ☐ Copy from a prior application (37 CFR 1.63(d))
(for continuation/divisional with Box 17 completed)
[Note Box 5 below]

12. ☒ Preliminary Amendment

i. ☐ DELETION OF INVENTOR(S)

Signed statement attached deleting inventor(s)
named in the prior application, see 37 CFR
1.63(d)(2) and 1.33(b).

13. ☒ Return Receipt Postcard (MPEP 503)
(Should be specifically itemized)

14. ☐ Small Entity Statement(s)

☐ Statement filed in prior application, Status still proper and desired

15. ☒ Certified Copy of Priority Document(s)

(if foreign priority is claimed)

16. ☐ Other

THE COMMISSIONER IS AUTHORIZED
TO CHARGE ANY DEFICIENCY IN THE
FEE FOR THIS PAPER TO DEPOSIT
ACCOUNT NO. 23-0975.

5. ☐ Incorporation By Reference
(usable if Box 4b is checked)

The entire disclosure of the prior application, from which
a copy of the oath or declaration is supplied under Box
4b, is considered as being part of the disclosure of the
accompanying application and is hereby incorporated by
reference therein.

17. If a CONTINUING APPLICATION, check appropriate box and supply the requisite information:

☐ Continuation ☐ Divisional ☐ Continuation-in-part (CIP) of prior Application No.

18. CORRESPONDENCE ADDRESS

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November 17, 2000

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of :
Yoichi MORI et al. :
Serial No. NEW : Attn: APPLICATION BRANCH
Filed November 17, 2000 : Attorney Docket No. 2000-1578A
METHOD AND APPARATUS FOR TREATING A WASTE GAS CONTAINING
FLUORINE-CONTAINING COMPOUNDS : THE COMMISSIONER IS AUTHORIZED
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FEE FOR THIS PAPER TO DEPOSIT
ACCOUNT NO. 23-0975.

PATENT OFFICE FEE TRANSMITTAL FORM

Assistant Commissioner for Patents
Washington, DC 20231

Sir:

Attached hereto is a check in the amount of \$750.00 to cover Patent Office fees relating to filing the following attached papers:

New application \$710.00

Assignment for Recordal \$ 40.00

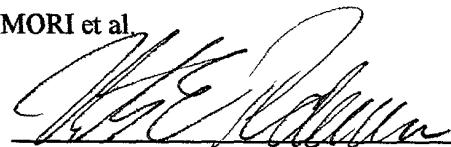
A duplicate copy of this paper is being submitted for use in the Accounting Division, Office of Finance.

The Commissioner is authorized to charge any deficiency or to credit any overpayment associated with this communication to Deposit Account No. 23-0975, with the EXCEPTION of deficiencies in fees for multiple dependent claims in new applications.

Respectfully submitted,

Yoichi MORI et al.

By:



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[Check No. 40799]
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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of :
Yoichi MORI et al. :
Serial No. NEW : **Attn: Application Branch**
Filed November 17, 2000 : **Attorney Docket No. 2000-1578A**

METHOD AND APPARATUS FOR
TREATING A WASTE GAS CONTAINING
FLUORINE-CONTAINING COMPOUNDS :

THE COMMISSIONER IS AUTHORIZED
TO CHARGE ANY DEFICIENCY IN THE
FEE FOR THIS PAPER TO DEPOSIT
ACCOUNT NO. 23-0975.

PRELIMINARY AMENDMENT

Assistant Commissioner for Patents
Washington, DC 20231

Sir:

Prior to examination of the instant application, please enter and consider the following
amendments and remarks as follows:

In The Claims:

Kindly amend claim 7 as follows:

Claim 7. line 1, delete "or 6".

Kindly add the following new claim 8:

-- 8. The apparatus according to claim 6, which has not only an air ejector capable
of adjusting the pressure in the apparatus through which the waste gas passes but also an FT-IR
analyzer for controlling the emission density of the treated gas.--

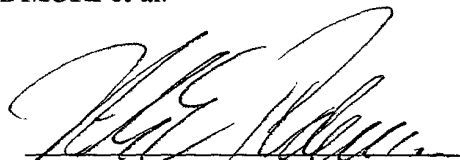
REMARKS

The present Preliminary Amendment is submitted to delete the multiple dependency of claim 8, thereby placing such claim in condition for examination and reducing the required PTO filing fee.

Respectfully submitted,

Yoichi MORI et al.

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METHOD AND APPARATUS FOR TREATING A WASTE GAS
CONTAINING FLUORINE-CONTAINING COMPOUNDS

BACKGROUND OF THE INVENTION

This invention relates to the treatment of a waste
5 gas containing fluorine-containing compounds. More
particularly, it relates to a method and an apparatus for
efficient treatment of emissions from semiconductor
fabrication plants, particularly from the steps of dry
cleaning the inner surface of a fabrication apparatus
10 and etching various kinds of deposited films with
perfluorocarbons (PFCs) and halogenated hydrocarbons such
as C_2F_6 , C_3F_8 , CHF_3 , SF_6 and NF_3 , said waste gases containing
not only PFCs but also oxidizing gases such as F_2 , Cl_2 and
 Br_2 , acidic gases such as HF , HCl , HBr , SiF_4 , $SiCl_4$, $SiBr_4$
15 and COF_2 , as well as CO .

Semiconductor fabrication plants use many kinds of
harmful gases that can potentially pollute the environment.
PFCs contained in waste gases that typically result from
the etching and CVD steps are suspected of causing global
20 warning and it is urgently needed to establish an effective
system for their removal.

Various breaking and recovery techniques have
heretofore been proposed for PFC removal. Catalytic
thermal decomposition is one of the breaking techniques and
25 uses versatile compounds such as Pt catalyst, zeolite-base
catalysts, activated charcoal, activated alumina, alkali
metals, alkaline earth metals and metal oxides. However,
none of these catalytic compounds have proved completely
satisfactory.

30 The waste gases discharged from the semiconductor
fabrication process contain not only PFCs but also
oxidizing gases such as F_2 , Cl_2 and Br_2 , acidic gases such
as HF , HCl , HBr , SiF_4 , $SiCl_4$, $SiBr_4$ and COF_2 , as well as CO ;
however, no method has yet been established that can
35 realize a thorough and effective treatment of these harmful
gases.

If one wants to treat oxidizing gases such as F_2 , Cl_2
and Br_2 by a wet method, thorough treatment cannot be

achieved by the sole use of water. If alkali agents or reducing agents are also used, not only process control but also the treatment apparatus becomes complicated and, in addition, the cost of treatment increases.

5 To remove CO, it has to be decomposed with oxidizers such as those based on Cu or Mn. As for PFCs, it has been proposed to use alumina as an agent for removing them (Japanese Patent Public Disclosure No. 286434/1998) and this method is characterized by contacting C₂F₆ with
10 molecular oxygen. However, the lifetime of alumina is very short and the throughput or the amount of C₂F₆ that can be treated for 100% decomposition is only 4.8 L/L; what is more, no effective way has been proposed to deal with CO that occurs as a by-product of C₂F₆ decomposition, still
15 less for the oxidizing gases and acidic gases that occur concomitantly with PFCs.

SUMMARY OF THE INVENTION

The present invention has been accomplished under these circumstances and has as an object providing a method
20 for treating waste gases containing fluorine-containing compounds which achieves high percent decomposition of PFCs, which proves effective for a prolonged time and which simultaneously realizes effective removal of oxidizing gases, acidic gases and CO that are contained in the waste
25 gases.

Another object of the invention is to provide an apparatus for implementing this method.

The first object of the invention can be attained by a method for treatment of a waste gas containing fluorine-
30 containing compounds which comprises the steps of separating the solids from the waste gas, adding H₂ and/or H₂O or H₂ and/or H₂O and O₂ as a decomposition assist gas, thermally decomposing the waste gas by contact with γ -alumina at usually at 500 - 1000°C, preferably 600 - 900°C, more
35 preferably 700 - 900°C, and removing acidic gases from the decomposed waste gas.

In this method, the waste gas containing fluorine-containing compounds may be a waste gas from a semiconductor

fabrication process that contains not only perfluorocarbons and fluorinated hydrocarbons but also oxidizing gases, acidic gases and CO.

The second object of the invention can be attained
5 by an apparatus for treatment of a waste gas containing
fluorine-containing compounds which comprises a solids
treating means for separating the solids from a waste gas
containing fluorine-containing compounds, an addition means
for adding H₂ and/or H₂O or H₂ and/or H₂O and O₂ as a
10 decomposition assist gas to the waste gas leaving the solids
treating means, a thermal decomposing means that is packed
with γ-alumina heated at 600 - 900°C and which thermally
decomposes the waste gas to which the decomposition assist
gas has been added, an acidic gas treating means for
15 removing acidic gases from the thermally decomposed waste
gas, and channels for connecting these means in sequence.

In this treatment apparatus, a water scrubber may be
used as the solids treating means or the acidic gas
treating means. This treatment apparatus may have not only
20 an air ejector capable of adjusting the pressure in the
apparatus through which the waste gas passes but also an
FT-IR analyzer for controlling the emission density of the
treated gas.

The first step in the method of the invention for
25 treating a waste gas containing fluorine-containing
compounds is passing the waste gas through a solids
treating means such as a water scrubber. The exit gas is
passed through a thermal decomposing means packed with
γ-alumina heated at 500 - 1000°C, preferably 600 - 900°C, and
30 more preferably 700 - 900°C, with H₂ and/or H₂O or H₂ and/or
H₂O and O₂ being added as a decomposition assist gas, so
that PFCs, oxidizing gases and CO are completely decomposed
into acidic gases and CO₂. The generated acidic gases are
removed with an acidic gas treating means such as a water
35 scrubber.

The invention may incorporate not only an air ejector
capable of adjusting the pressure in the apparatus through
which the waste gas passes but also an FT-IR analyzer for

controlling the emission density of the treated gas.

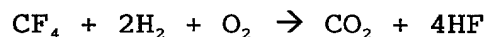
BRIEF DESCRIPTION OF THE DRAWING

Fig. 1 is a flowsheet for the waste gas treatment apparatus of the invention.

5 DETAILED DESCRIPTION OF THE INVENTION

The present invention is described below in detail. In the first step, a waste gas containing PFCs, oxidizing gases, acidic gases and CO is passed through a solids treating means such as a water scrubber in order to remove
10 not only the solids such as SiO_2 in the waste gas but also Si compounds such as SiF_4 , SiCl_4 and SiBr_4 that may potentially solidify in the thermal decomposing means at the next stage. If the waste gas is directly introduced into the thermal decomposing means without being passed
15 through the solids treating means, clogging or other blocking problems will occur, potentially preventing the waste gas from smoothly flowing through the packed γ -alumina layer. The performance of γ -alumina may also deteriorate. By passing the waste gas through the solids
20 treating means, the solids and the acidic gases containing Si compounds are removed whereas part of oxidizing gases such as F_2 , Cl_2 and Br_2 as well as all volumes of PFCs and CO are discharged.

The waste gas emerging from the solids treating means
25 is then introduced into the thermal decomposing means so that it is decomposed through contact with γ -alumina heated at $500 - 1000^\circ\text{C}$, preferably $600 - 900^\circ\text{C}$, more preferably $700 - 900^\circ\text{C}$. On this occasion, H_2 and/or H_2O ; or H_2 and/or H_2O and O_2 are added to the waste gas as a decomposition assist
30 gas so that the components of the waste gas are decomposed into acidic gases and CO_2 according to the following reaction schemes:



Thus, PFC reacts with H_2 and O_2 or H_2O to be decomposed into CO_2 and HF. Acidic gases such as F_2 react with H_2 or H_2O to be decomposed to another acidic gas HF. Carbon monoxide (CO) is oxidized to CO_2 .

5 As for PFC, H_2 or H_2O is added in moles at least equal to the moles necessary for the F atoms in the PFC to be converted to HF and O_2 is added in moles at least equal to the moles necessary for the C atoms in the PFC to be converted to CO_2 . Preferably, O_2 is added in moles which
10 are at least equal to the sum of one mole and the above-defined minimum number of moles. As for oxidizing gases, H_2 is introduced in moles at least equal to the moles necessary for the halogen atoms (X) in the oxidizing gas to be converted to an acidic gas (HX).

15 The waste gas leaving the thermal decomposing means only contains the acidic gases (HX) and CO_2 and by subsequent treatment with an acidic gas treating means such as a water scrubber, the acidic gases are completely removed.

20 The alumina to be used in the invention may have a γ -crystalline structure without a uniform pore distribution. While the shape of the alumina is not limited in any particular way, spheres are easy to handle and, hence, preferred. To the extent that will not unduly increase the
25 resistance to the passage of the waste gas, the particle size of γ -alumina should be as small as possible, preferably between 0.8 mm and 2.6 mm. The γ -alumina may be held at between $500^\circ C$ and $1000^\circ C$, preferably $600^\circ C$ and $900^\circ C$ and more preferably $700^\circ C$ and $900^\circ C$ during the passage of the waste
30 gas.

The solids treating means and the acidic gas treating means are preferably a packed column or a spray column on the condition that they are adapted to spray water. The thermal decomposing means should be adapted to permit the
35 introduction of H_2 and/or H_2O or H_2 and/or H_2O and O_2 as a decomposition assist gas.

Fig. 1 is a flowsheet for the waste gas treatment apparatus of the invention. It generally comprises the

solids treating means 1, the γ -alumina packed layer 2, the thermal decomposing means 3, a cleaning water circulating pump 4, the acidic gas treating means 5, an FT-IR analyzer 6, an air ejector 7 and a bypass valve 8.

5 A waste gas 9 containing PFCs, oxidizing gases, acidic gases and CO is first passed through the spray column (solids treating means) 1 so as to remove the solids and Si compounds. The waste gas is then passed through the thermal decomposing means 3 which is also supplied with H₂,
10 O₂ and H₂O to decompose the PFCs, oxidizing gases and CO into acidic gases and CO₂. The acidic gases are removed by passage through the next spray column (acidic gas treating means) 5, from which the treated gas 10 emerges.

The air ejector 7 is installed to control the pressure
15 in each of the treating means and the FT-IR analyzer 6 is provided to monitor the treated gas.

Spray water 11 is introduced into the acidic gas treating means 5 and the spent water is forced to the solids treating means 1 with the cleaning water circulating
20 pump 4; the water is used in spraying there and discharged as wastewater 12.

The following examples are provided for the purpose of further illustrating the present invention but are in no way to be taken as limiting.

25 Example 1

A quartz column of 25 mm ϕ was packed with γ -alumina to a height of 100 mm. The γ -alumina was a commercial product of Mizusawa Kagaku K.K. (NEOBEAD GB-08) having a particle size of 0.8 mm. The quartz column was installed in a
30 ceramic electric furnace and the alumina layer was heated at 800°C.

In addition to CF₄ diluted with N₂ gas, H₂ and O₂ were supplied as decomposition assist gases, the amount of H₂ being such that the number of H atoms was at least equal to
35 the number of F atoms in CF₄ and the amount of O₂ being at least equimolar to the amount of H₂ introduced. These gases were flowed into the column at a total rate of 408 sccm and their entrance concentrations were 1% (CF₄), 3.0%

(H₂) and 5.7% (O₂).

In order to evaluate the performance of the treatment system, the exit gas was analyzed periodically and the passage of the CF₄ gas was stopped when the removal of CF₄ dropped below 98%. The throughput was determined from the amount of CF₄ that had been passed through the system. The analysis of CF₄ and other gases was conducted with a gas chromatographic apparatus equipped with a mass detector.

As it turned out, the removal of CF₄ dropped to 98% when its passage continued for 920 min. At this point in time, the throughput as determined from the quantity of the supplied CF₄ was 77 L/L. Throughout the experiment, the concentration of CO emission was below the tolerable level (25 ppm).

15 Comparative Example 1

An experiment was conducted using the same equipment as in Example 1 that was packed with the same γ -alumina in the same amount and heated to the same temperature. The total gas flow rate was 408 sccm; the feed gas was a mixture of N₂-diluted CF₄ and SiF₄; in addition, H₂ and O₂ were supplied as decomposition assist gases, the amount of H₂ being such that the number of H atoms was at least equal to the total number of F atoms in CF₄ and SiF₄ and the amount of O₂ being at least equimolar to the amount of H₂ introduced. These gases were flowed into the column at respective concentrations of 0.95% (CF₄), 0.97% (SiF₄), 5.3% (H₂) and 6.0% (O₂).

As it turned out, the removal of CF₄ dropped below 98% when the passage of the CF₄/SiF₄ gas continued for 510 minutes. At this point in time, the throughput was 40 L/L, which was nearly one half the throughput for the case where only CF₄ gas was supplied. Throughout the experiment, the concentration of CO was below the tolerable level.

Example 2

35 An experiment was conducted using the same equipment as in Example 1 that was packed with the same γ -alumina in the same amount and heated to the same temperature. The total gas flow rate was 408 sccm; the feed gas was a

mixture of N₂-diluted CF₄ and F₂; in addition, H₂ and O₂ were supplied as decomposition assist gases, the amount of H₂ being such that the number of H atoms was at least equal to the total number of F atoms in CF₄ and F₂ and the amount of O₂ being at least equimolar to the amount of H₂ introduced. These gases were flowed into the column at respective concentrations of 0.92% (CF₄), 1.1% (F₂), 5.0% (H₂) and 6.0% (O₂).

As it turned out, the removal of CF₄ dropped below 98% when the passage of the CF₄/F₂ gas continued for 25 hours. At this point in time, the throughput was 115 L/L, which was 1.51 times higher than the throughput for the case where only CF₄ gas was supplied. Throughout the experiment, the concentrations of CO and F₂ were below the tolerable levels (1 ppm for F₂), provided that F₂ had been decomposed into HF.

Reference Example 1

An experiment was conducted using the same equipment as in Example 1 that was packed with the same γ -alumina in the same amount and heated to the same temperature. The total gas flow rate was 408 sccm; in addition to N₂-diluted CO, O₂ was supplied in moles at least equal to the moles necessary for CO to be converted to CO₂; their respective entrance concentrations were 1.4% (CO) and 5.7% (O₂). Throughout the passage of the feed gas for 30 minutes, the concentration of CO was below the detection limit (2 ppm); all of CO had been oxidized to CO₂.

Comparative Example 2

An experiment was conducted using the same equipment as in Example 1 that was packed with the same γ -alumina in the same amount and heated to the same temperature. The total gas flow rate was 408 sccm; in addition to N₂-diluted CO, H₂O was supplied at a rate of 0.090 mL/min which was 22 times as much as CO; the entrance concentration of CO was 1.3%.

As it turned out, 1000 ppm of CO leaked as a result of 15-min passage of the feed gas. Obviously, the concentration of CO could not be reduced to below the

tolerable level (25 ppm) by the sole addition of H₂O.

Reference Example 2

An experiment was conducted using the same equipment as in Example 1 that was packed with the same γ -alumina in the same amount and heated to the same temperature. The total gas flow rate was 408 sccm; in addition to N₂-diluted CO, H₂O was supplied at a rate of 0.090 mL/min which was 18 times as much as CO and O₂ was supplied in moles at least equal to the moles necessary for CO to be converted to CO₂; the entrance concentrations of CO and O₂ were 1.5% and 3.4%, respectively.

As it turned out, the concentration of CO had been reduced to below the detection limit (2 ppm) after the passage of the feed gas for 3 hours. Obviously, CO was oxidized to CO₂ by addition of O₂.

Example 3

An experiment was conducted using the same equipment as in Example 1 that was packed with the same γ -alumina in the same amount and heated to 700°C. The total gas flow rate was 408 sccm; in addition to N₂-diluted CF₄, H₂O was supplied at a rate of 0.040 mL/min which was 14 times as much as CF₄ and O₂ was supplied in moles at least equal to the moles necessary for the C atom in CF₄ to be converted to CO₂; the entrance concentrations of CF₄ and O₂ were 0.89% and 3.0%, respectively.

As it turned out, the removal of CF₄ dropped below 98% when the passage of the feed gas continued for 23 hours. At this point in time, the throughput was 110 L/L, which was 1.4 times higher than the throughput of CF₄ treatment in the presence of added H₂ and O₂. Throughout the experiment, the concentration of CO was below the tolerable level.

Comparative Example 3

In order to evaluate the effectiveness of the wet process in treating oxidizing gases and acidic gases, a water cleaning column (210 mm^φ x 430 mm^H with a Raschig ring packed to a height of 170 mm) was supplied with a waste gas at a total rate of 60 L/min and spray water at a

rate of 3.5 L/min. The waste gas was prepared from F_2 , SiF_4 and Cl_2 which had entrance concentrations of 1100 ppm, 1600 ppm and 5100 ppm, respectively. At the exit of the column, F_2 , SiF_4 and Cl_2 were detected at concentrations of 11 ppm, < 1 ppm, and 3300 ppm, respectively. Obviously, SiF_4 was effectively treated but F_2 and Cl_2 leaked out.

Example 4

A water cleaning column (210 mm ϕ x 430 mm H with a Raschig ring packed to a height of 170 mm) was used as a solids treating means. This column was combined with a thermal decomposing means comprising a preheating chamber and a catalyst packed chamber and an acidic gas treating means which was the same as the water cleaning column. The exit gas leaving the acidic gas treating means was monitored with an FT-IR analyzer (Infinity 6000 of MATTSON) and the pressure in the experimental system was controlled with an air ejector of Daito Seisakusho K.K. The solids treating means and the acidic gas treating means were supplied with cleaning water at respective flow rates of 2 L/min and 4 L/min. The thermal decomposing means was supplied with air and pure water at respective flow rates of 10 L/min and 2.4 mL/min. The catalyst was 15 L of γ -alumina (NEOBEAD GB-08 of Mizusawa Kagaku K.K.)

A gas dryer (MD-70-72P of PERMAPUR) was installed ahead of the FT-IR analyzer for removing the moisture in the waste gas. Air was supplied into the air ejector at a rate of 30 L/min so that the pressure in the system was kept at a negative value of -0.5 kPa. A waste gas was introduced at a flow rate of 60 L/min and it was prepared from a N_2 base containing CF_4 , SiF_4 , F_2 and CO at respective concentrations of 0.5%, 0.3%, 0.3% and 0.3%. The waste gas was first passed through the solids treating means, then passed through the thermal decomposing means in the presence of added water and O_2 , with the catalyst bed being heated at 700°C. The waste gas was subsequently passed through the acidic gas treating means and the treated gas was continuously analyzed by FT-IR. After the passage of the waste gas for 10 hours, CO_2 was detected in an amount

of 6900 ppm but each of CF_4 , SiF_4 , HF and CO had been treated to below 1 ppm. No F_2 was detected by ion chromatographic analysis.

Example 5

5 A waste gas treatment was conducted with the same experimental setup under the same conditions as in Example 4, except that CF_4 was replaced by C_2F_6 and that the waste gas was prepared from a N_2 base containing C_2F_6 , SiF_4 , F_2 and CO at respective concentrations of 0.5%, 0.3%, 0.3% and 10 0.3%. The waste gas was passed through the solids treating means, the thermal decomposing means and the acidic gas treating means.

15 The treated gas emerging from the acidic gas treating means was continuously analyzed by FT-IR. After the passage of the waste gas for 10 hours, CO_2 was detected in an amount of 11000 ppm but each of C_2F_6 , SiF_4 , HF and CO had been treated to below 1 ppm. No F_2 was detected by ion chromatographic analysis.

20 According to the invention, harmful waste gases from the semiconductor fabrication process that contain PFCs, oxidizing gases, acidic gases and CO and which are a potential accelerator of global warming can be treated in such a way that high percent decomposition is maintained for a prolonged time.

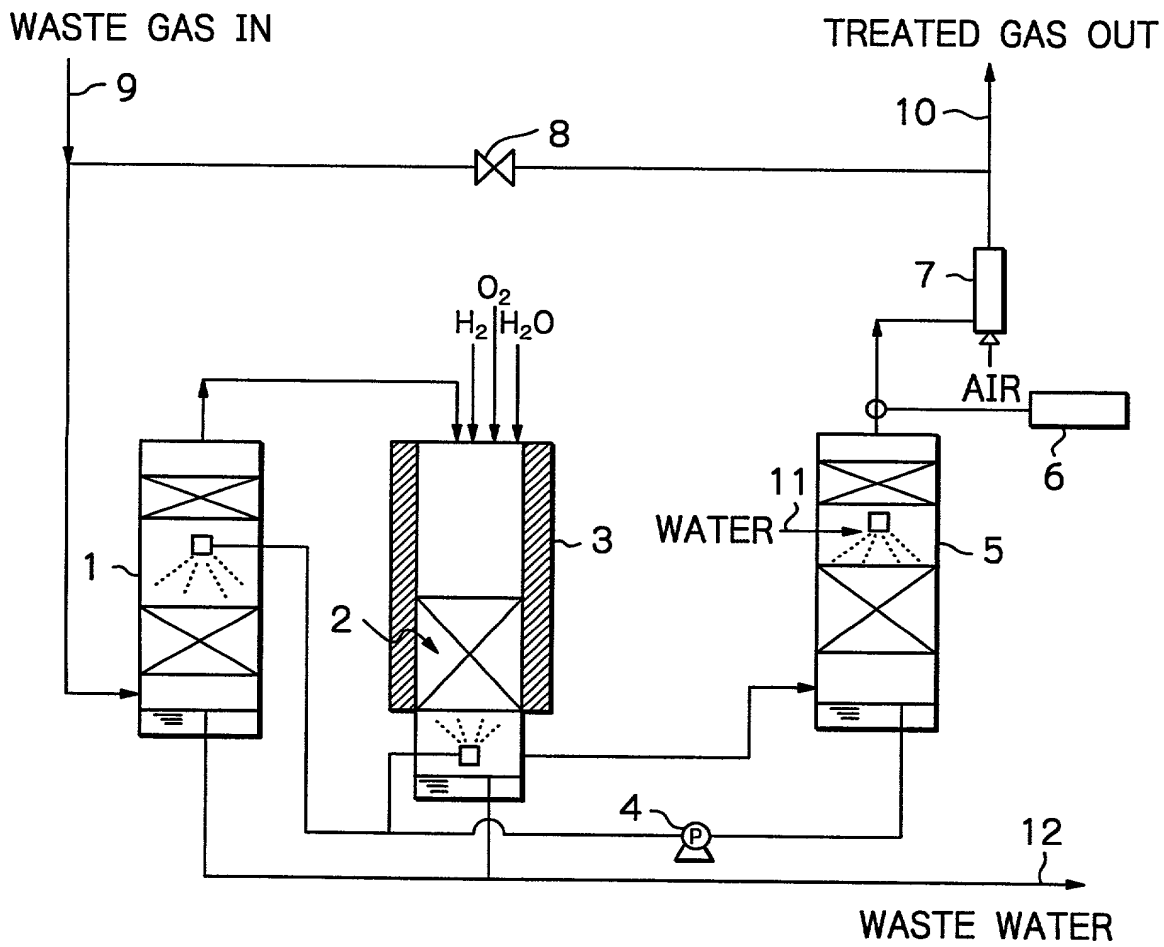
WHAT IS CLAIMED IS:

1. A method for treatment of a waste gas containing fluorine-containing compounds which comprises the steps of separating the solids from the waste gas, adding H_2 and/or H_2O or H_2 and/or H_2O and O_2 as a decomposition assist gas, thermally decomposing the waste gas by contact with γ -alumina at $500 - 1000^\circ C$, and removing acidic gases from the decomposed waste gas.
2. The method according to claim 1, wherein the temperature used in the thermally decomposing step ranges from $600 - 900^\circ C$
3. The method according to claim 1, wherein said waste gas containing fluorine-containing compounds contains perfluoro-carbons and fluorinated hydrocarbons as well as oxidizing gases, acidic gases and CO.
4. The method according to claim 3, wherein said waste gas containing fluorine-containing compounds is a waste gas from a semiconductor fabrication process.
5. An apparatus for treatment of a waste gas containing fluorine-containing compounds which comprises a solids treating means for separating the solids from a waste gas containing fluorine-containing compounds, an addition means for adding H_2 and/or H_2O ; or H_2 and/or H_2O and O_2 as a decomposition assist gas to the waste gas leaving the solids treating means, a thermal decomposing means that is packed with γ -alumina heated at $500 - 1000^\circ C$ and which thermally decomposes the waste gas to which the decomposition assist gas has been added, an acidic gas treating means for removing acidic gases from the thermally decomposed waste gas, and channels for connecting these means in sequence.
6. The apparatus according to claim 5, wherein said solids treating means or said acidic gas treating means is a water scrubber.
7. The apparatus according to claim 5 or 6, which has not only an air ejector capable of adjusting the pressure in the apparatus through which the waste gas passes but also an FT-IR analyzer for controlling the emission density of the treated gas.

ABSTRACT OF THE DISCLOSURE

The improved apparatus for treatment of a waste gas containing fluorine-containing compounds comprises solids treating means 1 for separating the solids from a waste gas containing fluorine-containing compounds, addition means for adding H_2 and/or H_2O or H_2 and/or H_2O and O_2 as a decomposition assist gas to the waste gas leaving the solids treating means, thermal decomposing means 3 that is packed with γ -alumina 2 heated at $600 - 900^\circ C$ and which thermally decomposes the waste gas to which the decomposition assist gas has been added, acidic gas treating means 5 for removing acidic gases from the thermally decomposed waste gas, and channels for connecting these means in sequence. The apparatus preferably includes air ejector 7 capable of adjusting its internal pressure.

Fig. 1



DECLARATION AND POWER OF ATTORNEY FOR U.S. PATENT APPLICATION

(X) Original () Supplemental () Substitute () PCT () Design

As a below named inventor, I hereby declare that: my residence, post office address and citizenship are as stated below next to my name; that I verily believe that I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural inventors are named below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

Title: METHOD AND APPARATUS FOR TREATING A WASTE GAS CONTAINING
FLUORINE-CONTAINING COMPOUNDS

of which is described and claimed in:

- (X) the attached specification, or
 () the specification in the application Serial No. _____ filed _____;
 and with amendments through _____ (if applicable), or
 () the specification in International Application No. PCT/ _____, filed _____, and as amended
 on _____ (if applicable).

I hereby state that I have reviewed and understand the content of the above-identified specification, including the claims, as amended by any amendment(s) referred to above.

I acknowledge my duty to disclose to the Patent and Trademark Office all information known to me to be material to patentability as defined in Title 37, Code of Federal Regulations, §1.56.

I hereby claim priority benefits under Title 35, United States Code, §119 (and §172 if this application is for a Design) of any application(s) for patent or inventor's certificate listed below and have also identified below any application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

COUNTRY	APPLICATION NO.	DATE OF FILING	PRIORITY CLAIMED
Japan	328411/1999	November 18, 1999	Yes

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose information material to patentability as defined in Title 37, Code of Federal Regulations, §1.56 which occurred between the filing date of the prior application and the national or PCT international filing date of this application.

APPLICATION SERIAL NO.	U.S. FILING DATE	STATUS: PATENTED, PENDING, ABANDONED

And I hereby appoint John T. Miller, Reg. No. 21,120; Michael R. Davis, Reg. No. 25,134; Matthew M. Jacob, Reg. No. 25,154; Jeffrey Nolton, Reg. No. 25,408; Warren M. Cheek, Jr., Reg. No. 33,367; Nils E. Pedersen, Reg. No. 33,145 and Charles R. Watts, Reg. No. 33,142, who together constitute the firm of WENDEROTH, LIND & PONACK, L.L.P., attorneys to prosecute this application and to transact all business in the U.S. Patent and Trademark Office connected therewith.

I hereby authorize the U.S. attorneys named herein to accept and follow instructions from YUASA AND HARA as to any action to be taken in the U.S. Patent and Trademark Office regarding this application without direct communication between the U.S. attorneys and myself. In the event of a change in the persons from whom instructions may be taken, the U.S. attorneys named herein will be so notified by me.

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I further declare that all statements made herein of my own knowledge are true, and that all statements on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

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The above application may be more particularly identified as follows:

U.S. Application Serial No. _____ Filing Date _____
 Applicant Reference Number _____ Atty Docket No. _____
 Title of Invention _____
